WO 2004/007513 PCT/KR2003/001398

What is claimed is:

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- 1. A method for producing 2-deoxy-L-ribose comprising the steps of
- (A) protection step for preparation of 2-deoxy-1-O-alkyl-D-ribopyranoside, of which the aldehyde group in 2-deoxy-D-ribose is protected in the form of acetal, by reacting 2-deoxy-D-ribose with an alcohol in the presence of an acid;
 - (B) activation step for preparation of 2-deoxy-1-O-alkyl-3,4-di-(alkanesulfonyl)-D-ribose or 2-deoxy-1-O-alkyl-3,4-di-(arylsulfonyl)-D-ribose, of which the 3-and 4-OH groups in 2-deoxy-D-ribose are activated, by reacting the above 2-deoxy-1-O-alkyl-D-ribose and an organic sulfonyl halide in the presence of a base;
 - (C) inversion step for preparation of a mixture of 2-deoxy-1-O-alkyl-3-acyl-L-ribose and 2-deoxy-1-O-alkyl-4-acyl-L-ribose, in which the stereochemistry of 3-OH and 4-OH groups are inverted, by reacting the above 2-deoxy-1-O-alkyl-3,4-di-(alkanesulfonyl)-D-ribose or 2-deoxy-1-O-alkyl-3,4-di-(arylsulfonyl)-D-ribose with a metal salt of organic acid; and
 - (D) deprotection step for preparation for 2-deoxy-L-ribose by reactions of the above mixture of 2-deoxy-1-O-alkyl-3-acyl-L-ribose and 2-deoxy-1-O-alkyl-4-acyl-L-ribose with an acid and then with a base, or with a base and then with an acid.
- 20 2. The method for producing 2-deoxy-L-ribose according to Claim 1, wherein the alcohol used in said protection step is a lower alcohol having 1-4 of carbon number, benzyl alcohol or substituted benzyl alcohol.
 - 3. The method for producing 2-deoxy-L-ribose according to Claim 1, wherein the

WO 2004/007513 PCT/KR2003/001398

organic sulfonyl halide used in said activation step is a lower alkanesulfonyl halide such as methane sulfonyl chloride or trifluoromethyl chloride, or arylsulfonyl halide such as benzenesulfonyl chloride or p-toluenesulfonyl chloride.

4. The method for producing 2-deoxy-L-ribose according to Claim 1, wherein the metal salt of organic acid used in said conversion step is a metal salt of lower alkyl organic acid having 1-8 of carbon number or a metal salt of aryl organic acid.